## **Supplementary Online Content**

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This supplementary material has been provided by the authors to give readers additional information about their work.

eTable 1. List of clinical trials

Trial Name	Phase	Treatment Arm	Time	N	PASI 50	PASI 75	PASI 90	PASI 100
			Point		(%)	(%)	(%)	(%)
Clinical trials incl	uded in the	short-term NMA						
Asahina	2/3	Placebo	16	46	19.6	4.4	0	
		Adalimumab 80 mg at week 0, then 40 mg						
Asahina	2/3	EOW starting at week 1	16	43	81.4	62.8	39.5	
Bissonnette	4	Placebo	16	10		20		
		Adalimumab 80 mg at week 0, then 40 mg						
Bissonnette	4	EOW starting at week 1	16	20		70		
REVEAL	3	Placebo	16	398		6.5		
		Adalimumab 80 mg at week 0, then 40 mg						
REVEAL	3	EOW starting at week 1	16	814		71		
CHAMPION	3	Placebo	16	53	30.2	18.9	11.3	1.9
CHAMPION	3	Methotrexate (low dose)	16	110	51.8	35.5	13.6	7.3
		Adalimumab 80 mg at week 0, then 40 mg						
CHAMPION	3	EOW starting at week 1	16	108	88	79.6	51.9	16.7
Gordon 2006	2	Placebo	12	52		3.9		0

Trial Name	Phase	Treatment Arm	Time	N	PASI 50	PASI 75	PASI 90	PASI 100
			Point		(%)	(%)	(%)	(%)
		Adalimumab 80 mg at week 0, then 40 mg						
Gordon 2006	2	EOW starting at week 1	12	45		53.3		11.1
Cai 2017	3	Placebo	12	87		11.5	3.4	1.1
		Adalimumab 80 mg at week 0, then 40 mg						
Cai 2017	3	EOW starting at week 1	12	338		77.8	55.6	13.3
Goldminz 2015	4	Methotrexate (low dose)	16	15		26.7		
		Adalimumab 80 mg at week 0, then 40 mg						
Goldminz 2015	4	EOW starting at week 1	16	15		66.7		
Leonardi	3	Placebo	12	166	14.5	3.6	0.6	
Leonardi	3	Etanercept 25 mg BIW / 50 mg QW	12	162	58	34	11.7	
Papp	3	Placebo	12	193	9.3	3.1	0.5	
Papp	3	Etanercept 25 mg BIW / 50 mg QW	12	196	64.3	34.2	10.7	
van de Kerkhof	3	Placebo	12	46	8.7	2.2	2.2	
van de Kerkhof	3	Etanercept 25 mg BIW / 50 mg QW	12	96	68.8	37.5	13.5	
Gottlieb	2	Placebo	12	55	10.9	1.8	0	
Gottlieb	2	Etanercept 25 mg BIW / 50 mg QW	12	57	70.2	29.8	10.5	
EXPRESS	3	Placebo	10	77	7.8	2.6	1.3	

Trial Name	Phase	Treatment Arm	Time	N	PASI 50	PASI 75	PASI 90	PASI 100
			Point		(%)	(%)	(%)	(%)
		Infliximab 5 mg/kg at weeks 0, 2 and 6, then						
EXPRESS	3	Q8W	10	301	91	80.4	57.1	
EXPRESS II	3	Placebo	10	208		1.9	0.5	
		Infliximab 5 mg/kg at weeks 0, 2 and 6, then						
EXPRESS II	3	Q8W	10	314		75.5	45.2	
SPIRIT	2	Placebo	10	51	21.6	5.9	2	
		Infliximab 5 mg/kg at weeks 0, 2 and 6, then						
SPIRIT	2	Q8W	10	99	97	87.9	57.6	
Chaudhari		Placebo	10	11		18.2		
		Infliximab 5 mg/kg at weeks 0, 2 and 6, then						
Chaudhari		Q8W	10	11		81.8		
Torii	3	Placebo	10	19	10.5	0	0	
		Infliximab 5 mg/kg at weeks 0, 2 and 6, then						
Torii	3	Q8W	10	35	82.9	68.6	54.3	
Yang		Placebo	10	45	13.3	2.2	0	
		Infliximab 5 mg/kg at weeks 0, 2 and 6, then						
Yang		Q8W	10	84	94.1	81	57.1	
UNCOVER 1	3	Placebo	12	431	11.6	3.9	0.5	0

Trial Name	Phase	Treatment Arm	Time	N	PASI 50	PASI 75	PASI 90	PASI 100
			Point		(%)	(%)	(%)	(%)
		Ixekizumab 160 mg at week 0, then 80 mg						
UNCOVER 1	3	Q2W	12	433	93.8	89.2	70.9	35.3
UNCOVER 2	3	Placebo	12	168	6.6	2.4	0.6	0.6
		Ixekizumab 160 mg at week 0, then 80 mg						
UNCOVER 2	3	Q2W	12	351	94.9	89.7	70.7	40.5
UNCOVER 3	3	Placebo	12	193	15.5	7.3	3.1	0
		Ixekizumab 160 mg at week 0, then 80 mg						
UNCOVER 3	3	Q2W	12	385	93.8	87.3	68.1	37.7
		Ixekizumab 160 mg at week 0, then 80 mg						
IXORA-S	3	Q2W	12	136		88.2	72.8	36
		Ustekinumab 45 mg < 100 kg, 90 mg > 100 kg						
IXORA-S	3	at weeks 0 and 4, then Q12W	12	166		68.7	42.2	14.5
		Ixekizumab 160 mg at week 0, then 80 mg						
Reich 2017	3	Q2W	12	54		90.7	72.2	35.2
Reich 2017	3	Methotrexate (low dose)	12	54		48.1	20.4	5.6
ERASURE	3	Placebo	12	246	8.9	4.5	1.2	0.8
		Secukinumab 300 mg at weeks 0, 1, 2, and 3,						
ERASURE	3	then monthly starting at week 4	12	245	90.6	81.6	59.2	28.6

Trial Name	Phase	Treatment Arm	Time	N	PASI 50	PASI 75	PASI 90	PASI 100
			Point		(%)	(%)	(%)	(%)
FEATURE	3	Placebo	12	59	5.1	0	0	0
		Secukinumab 300 mg at weeks 0, 1, 2, and 3,						
FEATURE	3	then monthly starting at week 4	12	58	87.9	75.9	60.3	43.1
FIXTURE	3	Placebo	12	324	15.1	4.9	1.5	0
		Secukinumab 300 mg at weeks 0, 1, 2, and 3,						
FIXTURE	3	then monthly starting at week 4	12	323	91.6	77.1	54.2	24.2
JUNCTURE	3	Placebo	12	61	8.2	3.3	0	0
		Secukinumab 300 mg at weeks 0, 1, 2, and 3,						
JUNCTURE	3	then monthly starting at week 4	12	60	96.7	86.7	55	26.7
		Ustekinumab 45 mg < 100 kg, 90 mg > 100 kg						
CLEAR	3	at weeks 0 and 4, then Q12W	12	335		79.1	53.4	25.7
		Secukinumab 300 mg at weeks 0, 1, 2, and 3,						
CLEAR	3	then monthly starting at week 4	12	334		91	72.8	38.9
PRIME	3	Fumaric acid esters	12	95	56.8	21.1	2.1	0
		Secukinumab 300 mg at weeks 0, 1, 2, and 3,						
PRIME	3	then monthly starting at week 4	12	105	97.1	87.6	63.8	28.6
		Ustekinumab 45 mg < 100 kg, 90 mg > 100 kg						
CLARITY	3	at weeks 0 and 4, then Q12W	12	552		74.3	47.8	20.1

Trial Name	Phase	Treatment Arm	Time	N	PASI 50	PASI 75	PASI 90	PASI 100
			Point		(%)	(%)	(%)	(%)
		Secukinumab 300 mg at weeks 0, 1, 2, and 3,						
CLARITY	3	then monthly starting at week 4	12	550		88	66.5	38.2
		Ustekinumab 45 mg at weeks 0 and 4, then						
ACCEPT	3	Q12W	12	209		67.5	36.4	
		Ustekinumab 90 mg at weeks 0 and 4, then						
ACCEPT	3	Q12W	12	347		73.8	44.7	
LOTUS	3	Placebo	12	162	19.8	11.1	3.1	0.6
		Ustekinumab 45 mg at weeks 0 and 4, then						
LOTUS	3	Q12W	12	160	91.3	82.5	66.9	23.8
PEARL	3	Placebo	12	60	13.3	5	1.7	0
		Ustekinumab 45 mg at weeks 0 and 4, then						
PEARL	3	Q12W	12	61	83.6	67.2	49.2	8.2
PHOENIX 1	3	Placebo	12	255	10.2	3.1	2	0
		Ustekinumab 45 mg at weeks 0 and 4, then						
PHOENIX 1	3	Q12W	12	255	83.5	67.1	41.6	12.6
		Ustekinumab 90 mg at weeks 0 and 4, then						
PHOENIX 1	3	Q12W	12	256	85.9	66.4	36.7	10.9
PHOENIX 2	3	Placebo	12	410	10	3.7	0.7	0

Trial Name	Phase	Treatment Arm	Time	N	PASI 50	PASI 75	PASI 90	PASI 100
			Point		(%)	(%)	(%)	(%)
		Ustekinumab 45 mg at weeks 0 and 4, then						
PHOENIX 2	3	Q12W	12	409	83.6	66.8	42.3	18.1
		Ustekinumab 90 mg at weeks 0 and 4, then						
PHOENIX 2	3	Q12W	12	411	89.3	75.7	50.9	18.3
Igarashi	2/3	Placebo	12	31	12.9	6.5	3.2	
		Ustekinumab 45 mg at weeks 0 and 4, then						
Igarashi	2/3	Q12W	12	64	82.8	59.4	32.8	
		Ustekinumab 90 mg at weeks 0 and 4, then						
Igarashi	2/3	Q12W	12	62	83.9	67.7	43.6	
VIP-U	4	Placebo	12	21		9.5		
		Ustekinumab 45 mg < 100 kg, 90 mg > 100 kg						
VIP-U	4	at weeks 0 and 4, then Q12W	12	22		77.3		
X-PLORE	2	Placebo	16	42		4.8	2.4	0
		Adalimumab 80 mg at week 0, then 40 mg						
X-PLORE	2	EOW starting at week 1	16	43		69.8	44.2	25.6
VOYAGE-1	3	Placebo	16	174		5.7	2.9	0.6
		Adalimumab 80 mg at week 0, then 40 mg						
VOYAGE-1	3	EOW starting at week 1	16	334		73.1	49.7	17.1

Trial Name	Phase	Treatment Arm	Time	N	PASI 50	PASI 75	PASI 90	PASI 100
			Point		(%)	(%)	(%)	(%)
VOYAGE-1	3	Guselkumab 100 mg at weeks 0, 4, then Q8W	16	329		91.2	73.3	37.4
VOYAGE-2	3	Placebo	16	248		8.1	2.4	0.8
		Adalimumab 80 mg at week 0, then 40 mg						
VOYAGE-2	3	EOW starting at week 1	16	248		68.5	46.8	20.6
VOYAGE-2	3	Guselkumab 100 mg at weeks 0,4, then Q8W	16	496		86.3	70	34.1
ORION	3	Placebo	16	16			0	0
ORION	3	Guselkumab 100 mg at weeks 0, 4, then Q8W	16	62			75.8	50
Ohtsuki 2018	3	Placebo	16	64	14.1	6.3	0	0
Ohtsuki 2018	3	Guselkumab 100 mg at weeks 0, 4, then Q8W	16	63	95.2	84.1	69.8	27
Nakagawa 2016	2	Placebo	12	38		7.9	2.6	0
		Brodalumab 210 mg at weeks 0, 1, and 2, then						
Nakagawa 2016	2	Q2W	12	37		94.6	91.9	59.5
Papp 2012	2	Placebo	12	38	15.8	0	0	0
		Brodalumab 210 mg at weeks 0, 1, and 2, then						
Papp 2012	2	Q2W	12	40	90	82.5	75	62.5
AMAGINE-1	3	Placebo	12	220		2.7	0.9	0.5
		Brodalumab 210 mg at weeks 0, 1, and 2, then						
AMAGINE-1	3	Q2W	12	222		83.3	70.3	41.9

Trial Name	Phase	Treatment Arm	Time	N	PASI 50	PASI 75	PASI 90	PASI 100
			Point		(%)	(%)	(%)	(%)
AMAGINE-2	3	Placebo	12	309		8.1	2.9	0.7
		Ustekinumab 45 mg < 100 kg, 90 mg > 100 kg						
AMAGINE-2	3	at weeks 0 and 4, then Q12W	12	300		70	47	21.7
		Brodalumab 210 mg at weeks 0, 1, and 2, then						
AMAGINE-2	3	Q2W	12	612		86.3	70	44.4
AMAGINE-3	3	Placebo	12	315		6	1.9	0.3
		Ustekinumab 45 mg < 100 kg, 90 mg > 100 kg						
AMAGINE-3	3	at weeks 0 and 4, then Q12W	12	313		69.3	47.9	18.5
		Brodalumab 210 mg at weeks 0, 1, and 2, then						
AMAGINE-3	3	Q2W	12	624		85.1	69	36.7
CIMPASI-1	3	Placebo	16	51		6.5	0.4	0.2
		Certolizumab pegol 400 mg at weeks 0, 2, and						
CIMPASI-1	3	4, then 200 mg Q2W	16	95		66.5	35.8	13.7
CIMPASI-1	3	Certolizumab pegol 400 mg Q2W	16	88		75.8	43.6	12.7
CIMPASI-2	3	Placebo	16	49		11.6	4.5	1.8
		Certolizumab pegol 400 mg at weeks 0, 2, and						
CIMPASI-2	3	4, then 200 mg Q2W	16	91		81.4	52.6	15.4
CIMPASI-2	3	Certolizumab pegol 400 mg Q2W	16	87		82.6	55.4	18.8

Trial Name	Phase	Treatment Arm	Time	N	PASI 50	PASI 75	PASI 90	PASI 100
			Point		(%)	(%)	(%)	(%)
CIMPACT	2	Placebo	12	57		5	0.2	
		Certolizumab pegol 400 mg at weeks 0, 2, and						
CIMPACT	2	4, then 200 mg Q2W	12	165		61.3	31.2	
CIMPACT	2	Certolizumab pegol 400 mg Q2W	12	167		66.7	34	
NCT00245765	2	Placebo	12	59	11.9	6.8	1.7	
		Certolizumab pegol 400 mg at week 0, then						
NCT00245765	2	200 mg Q2W	12	59	86.4	74.6	39	
NCT00245765	2	Certolizumab pegol 400 mg Q2W	12	58	93.1	82.8	46.6	
reSURFACE-1	3	Placebo	12	154		5.8	2.6	1.3
		Tildrakizumab 100 mg at weeks 0, 4, then						
reSURFACE-1	3	Q12W	12	309		63.8	34.6	13.9
		Tildrakizumab 200 mg at weeks 0, 4, then						
reSURFACE-1	3	Q12W	12	308		62.3	35.4	14
Papp 2015	2	Placebo	16	45		4.4	2.2	
		Tildrakizumab 100 mg at weeks 0, 4, then						
Papp 2015	2	Q12W	16	89		66.3	38.2	
		Tildrakizumab 200 mg at weeks 0, 4, then						
Papp 2015	2	Q12W	16	86		74.4	51.2	

Trial Name	Phase	Treatment Arm	Time	N	PASI 50	PASI 75	PASI 90	PASI 100
			Point		(%)	(%)	(%)	(%)
reSURFACE-2	3	Placebo	12	156		6	1	0
		Tildrakizumab 100 mg at weeks 0, 4, then						
reSURFACE-2	3	Q12W	12	307		61	39	12.4
		Tildrakizumab 200 mg at weeks 0, 4, then						
reSURFACE-2	3	Q12W	12	314		66	37	11.8
UltIMMa1	3	Placebo	16	102	21.6	8.8	4.9	0
		Ustekinumab 45 mg < 100 kg, 90 mg > 100 kg						
UltIMMa1	3	at weeks 0 and 4, then Q12W	16	100	89	76	42	12
		Risankizumab 150 mg at weeks 0, and 4, then						
UltIMMa1	3	Q12W	16	304	94.1	89.1	75.3	35.9
UltIMMa2	3	Placebo	16	98	16.3	6.1	2	2
		Ustekinumab 45 mg < 100 kg, 90 mg > 100 kg						
UltIMMa2	3	at weeks 0 and 4, then Q12W	16	99	80.8	69.7	47.5	24.2
		Risankizumab 150 mg at weeks 0, and 4, then						
UltIMMa2	3	Q12W	16	294	97.6	90.8	74.8	50.7
		Adalimumab 80 mg at week 0, then 40 mg						
IMMvent	3	EOW starting at week 1	16	304	83.6	71.7	47.4	23

Trial Name	Phase	Treatment Arm	Time	N	PASI 50	PASI 75	PASI 90	PASI 100
			Point		(%)	(%)	(%)	(%)
		Risankizumab 150 mg at weeks 0, and 4, then						
IMMvent	3	Q12W	16	301	97	90.7	72.4	39.9
IMMhance	3	Placebo	16	100	19	8	2	1
		Risankizumab 150 mg at weeks 0, and 4, then						
IMMhance	3	Q12W	16	407	95.6	88.7	73.2	47.2
BRIDGE	3	Placebo	16	131	29	15.3	4.6	
BRIDGE	3	Fumaric acid esters	16	273	61.9	40.3	22.3	
BRIDGE	3	Dimethyl fumarate	16	267	53.6	37.5	18.4	
Altmeyer 1994		Placebo	16	51		2		
Altmeyer 1994		Fumaric acid esters	16	49		24.5		
PSOR-008 /								
ESTEEM-1	3	Placebo	16	282	17	5.3	0.4	
PSOR-008 /		Apremilast 30 mg twice daily after initial						
ESTEEM-1	3	titration schedule	16	562	58.7	33.1	9.8	
PSOR-009 /								
ESTEEM-2	3	Placebo	16	137	19.7	5.8	1.5	
PSOR-009 /		Apremilast 30 mg twice daily after initial						
ESTEEM-2	3	titration schedule	16	274	55.5	28.8	8.8	

Trial Name	Phase	Treatment Arm	Time	N	PASI 50	PASI 75	PASI 90	PASI 100
			Point		(%)	(%)	(%)	(%)
PSOR-010 /								
LIBERATE	3b	Placebo	16	84	33.3	11.9	3.6	
PSOR-010 /								
LIBERATE	3b	Etanercept 25 mg BIW / 50 mg QW	16	83	83.1	48.2	20.5	
PSOR-010 /		Apremilast 30 mg twice daily after initial						
LIBERATE	3b	titration schedule	16	83	62.7	39.8	14.5	
PSOR-005	2b	Placebo	16	88	25	5.7	1.1	
		Apremilast 30 mg twice daily after initial						
PSOR-005	2b	titration schedule	16	88	60.2	40.9	11.4	
Ohtsuki 2017	2b	Placebo	16	84	21.4	7.1	1.2	
		Apremilast 30 mg twice daily after initial						
Ohtsuki 2017	2b	titration schedule	16	85	50.6	28.2	14.1	
Gisondi 2008		Etanercept 25 mg BIW / 50 mg QW	12	22	40.9	22.7		
Gisondi 2008		Acitretin 0.4 mg/kg daily	12	20	20	10		
Meffert 1997		Placebo	10	43	11.6	4.7		
Meffert 1997		Ciclosporin 2.5–3 mg/kg/day	10	44	56.8	29.5		
		Infliximab 5 mg/kg at weeks 0, 2 and 6, then						
RESTORE1	3b	Q8W	16	653	86.8	77.8	54.5	

Trial Name	Treatment Arm		Time	N	PASI 50	PASI 75	PASI 90	PASI 100
			Point		(%)	(%)	(%)	(%)
RESTORE1	3b	Methotrexate (high dose)	16	215	60.5	41.9	19.1	
Heydendael 2003		Methotrexate (high dose)	16	43		60.5	39.5	
Heydendael 2003		Ciclosporin 2.5–3 mg/kg/day	16	42		71.4	33.3	
Fallah 2011		Methotrexate (low dose)	12	25	60	24	8	
Fallah 2011		Fumaric acid esters	12	26	42.3	19.2	3.8	
Flytstrom 2007		Methotrexate (low dose)	12	37	64.9	24.3	10.8	
Flytstrom 2007		Ciclosporin 2.5–3 mg/kg/day	12	31	87.1	58.1	29	
Clinical trials inclu	ided in the	e long-term meta-analysis	l					
LIBERATE	3b	Apremilast 30 mg BID	52	74	70.3	52.7	17.6	
Ohtsuki 2017	2b	Apremilast 30 mg BID	52	85		40.0		
PSOR-005/PSOR-	2	Apremilast 30 mg BID	52	58	72.4	36.2	13.8	
005E								
FIXTURE	3	Etanercept 50 mg BIW until week 12, then	52	323		55.5	33.4	10.1
		QW						
EXPRESS	3	Infliximab 5 mg/kg at week 0, 2, and 6, then	50	281	68.7	60.5	45.2	
		Q8W						

Trial Name	Phase	Treatment Arm	Time	N	PASI 50	PASI 75	PASI 90	PASI 100
			Point		(%)	(%)	(%)	(%)
EXPRESS II	3	Infliximab 5 mg/kg at week 0, 2, and 6, then	50	134	72.4	54.5	34.3	
		Q8W						
ADACCESS	3	Adalimumab 80 mg at week 0, then 40 mg	51	115	93.9	79.6	51.0	29.6
		Q2W						
Gordon 2006	2	Adalimumab 80 mg at week 0, then 40 mg	60	45	64.0	56.0	33.0	16.0
		Q2W						
VOYAGE 1	3	Adalimumab 80 mg at week 0, then 40 mg	48	334		62.6	47.9	23.4
		Q2W						
AMAGINE-2	3	Ustekinumab 45 mg or 90 mg at week 0, 4,	52	300		62.0	48.0	30.0
		then Q12W						
AMAGINE-3	3	Ustekinumab 45 mg or 90 mg at week 0, 4,	52	313		63.0	50.0	29.0
		then Q12W						
CLEAR	3b	Ustekinumab 45 mg or 90 mg at week 0, 4,	52	335		78.2	60.6	36.7
		then Q12W						
IXORA-S	3	Ustekinumab 45 mg or 90 mg at week 0, 4,	52	166		75.9	59.0	35.5
		then Q12W						
PSTELLAR	3	Ustekinumab 45 mg or 90 mg at week 0, 4,	52	69		82.6		
		then Q12W						

Trial Name	Phase Treatment Arm			N	PASI 50	PASI 75	PASI 90	PASI 100
			Point		(%)	(%)	(%)	(%)
UltIMMa1	3	Ustekinumab 45 mg or 90 mg at week 0, 4,	52	100	87.0	70.0	44.0	21.0
		then Q12W						
UltIMMa2	3	Ustekinumab 45 mg or 90 mg at week 0, 4,	52	99	85.9	76.8	50.5	30.3
		then Q12W						
CLEAR	3b	Secukinumab 300 mg at week 0, 1, 2, and 3,	52	334		91.6	74.9	44.9
		then Q4W						
Pooled analysis of	3	Secukinumab 300 mg at week 0, 1, 2, and 3,	52	686		85.2	68.1	40.8
four trials		then Q4W						
(ERASURE,								
FEATURE,								
FIXTURE, and								
JUNCTURE)								
IXORA-S	3	Ixekizumab 160 mg at week 0, 80 mg Q2W	52	136		88.2	76.5	52.2
		until week 12, then 80 mg Q4W						
UNCOVER 3	3	Ixekizumab 160 mg at week 0, 80 mg Q2W	60	385		83.0	73.0	55.0
		until week 12, then 80 mg Q4W						
AMAGINE-2	3	Brodalumab 210 mg at week 0, 1, 2, then Q2W	52	189		80.0	75.0	56.0
AMAGINE-3	3	Brodalumab 210 mg at week 0, 1, 2, then Q2W	52	194		80.0	73.0	53.0

Trial Name	Phase	Treatment Arm	Time	N	PASI 50	PASI 75	PASI 90	PASI 100
			Point		(%)	(%)	(%)	(%)
Ohtsuki 2018	3	Guselkumab 100 mg at week 0, 4, then Q8W	52	63	98.4	90.5	77.8	47.6
VOYAGE 1	3	Guselkumab 100 mg at week 0, 4, then Q8W	48	329		87.8	76.3	47.4
IMMvent	3	Risankizumab 150 mg at week 0, 4, then Q12W	44	301	89.0	86.7	75.7	52.8
UltIMMa1	3	Risankizumab 150 mg at week 0, 4, then Q12W	52	304	94.4	91.8	81.9	56.3
UltIMMa2	3	Risankizumab 150 mg at week 0, 4, then Q12W	52	294	93.2	91.5	80.6	59.5

BID, twice daily; BIW, twice a week; EOW, every other week; kg, kilogram; mg, milligram; NMA, network-meta analysis; PASI, Psoriasis Area and Severity Index; Q12W, once every 12 weeks; Q2W, once every 2 weeks; Q4W, once every 4 weeks; Q8W, once every 8 weeks; QW, once weekly

**eFigure 1.** Estimated odds ratios from the NMA of short-term PASI (base-case)

	Risankizumab 150 mg		lxekizumab 80 mg		Brodalumab 210 mg		Guselkumab 100 mg	
Risankizumab 150 mg PASI 75 PASI 90 PASI 100		-	H	0.96 (0.70, 1.31) 0.96 (0.73, 1.26) 0.96 (0.74, 1.25)	H-H-H-H-H-H-H-H-H-H-H-H-H-H-H-H-H-H-H-	0.95 (0.71, 1.29) 0.95 (0.74, 1.24) 0.95 (0.75, 1.24)		0.79 (0.58, 1.08) 0.81 (0.62, 1.07) 0.82 (0.63, 1.07)
bxekizumab 80 mg PASI 75 PASI 90 PASI 100		1.05 (0.76, 1.43) 1.04 (0.79, 1.36) 1.04 (0.80, 1.35)		-		0.99 (0.74, 1.36) 0.99 (0.77, 1.30) 0.99 (0.77, 1.29)	He-H	0.83 (0.60, 1.15) 0.85 (0.64, 1.13) 0.85 (0.64, 1.13)
Brodalumab 210 mg PASI 75 PASI 90 PASI 100	H-H	1.05 (0.80, 1.34)	 	1.01 (0.74, 1.35) 1.01 (0.77, 1.30) 1.01 (0.77, 1.29)		:	H-H-H-H-H-H-H-H-H-H-H-H-H-H-H-H-H-H-H-	0.84 (0.61, 1.13) 0.85 (0.65, 1.12) 0.86 (0.65, 1.11)
Guselkumab 100 mg PASI 75 PASI 90 PASI 100	H- <b>6</b> -1 H-6-1 H-6-1	1.27 (0.93, 1.72) 1.23 (0.93, 1.60) 1.22 (0.94, 1.59)	 	1.21 (0.87, 1.68) 1.18 (0.88, 1.57) 1.18 (0.89, 1.56)	H	1.20 (0.88, 1.65) 1.17 (0.90, 1.55) 1.17 (0.90, 1.53)		-
Secukinumab 300 mg PASI 75 PASI 90 PASI 100	<b>1-8-1</b>   1-8-1   1-8-1	1.69 (1.26, 2.26)* 1.59 (1.23, 2.05)* 1.58 (1.23, 2.04)*	H	1.62 (1.22, 2.14)* 1.53 (1.19, 1.96)* 1.53 (1.19, 1.95)*	   <del>                                   </del>	1.60 (1.22, 2.12)* 1.51 (1.20, 1.94)* 1.51 (1.20, 1.94)*		1.34 (0.98, 1.83) 1.29 (0.98, 1.71) 1.30 (0.98, 1.71)
Infliximab 5 mg/kg PASI 75 PASI 90 PASI 100	<b></b>	2.03 (1.44, 2.83)* 1.87 (1.38, 2.52)* 1.88 (1.38, 2.54)*	   H#H   H#H	1.94 (1.41, 2.66)* 1.80 (1.35, 2.39)* 1.81 (1.36, 2.41)*	<b></b>	1.92 (1.39, 2.68)* 1.78 (1.34, 2.41)* 1.79 (1.34, 2.43)*	 	1.60 (1.12, 2.27)* 1.52 (1.11, 2.09)* 1.54 (1.11, 2.12)*
Certolizumab pegol 400 mg PASI 75 PASI 90 PASI 100	<b>⊢●</b> -1 <b>⊢●</b> -1 <b>⊢●</b> -1	3.37 (2.37, 4.76)* 3.02 (2.18, 4.15)* 3.15 (2.24, 4.44)*	H=-1 H=-1	3.22 (2.28, 4.56)* 2.90 (2.11, 4.00)* 3.03 (2.16, 4.29)*	    	3.19 (2.27, 4.52)* 2.88 (2.10, 3.97)* 3.01 (2.15, 4.25)*	 	2.67 (1.85, 3.81)* 2.46 (1.76, 3.43)* 2.58 (1.81, 3.68)*
Ustekinumab 45 mg ≤100 kg, 90 mg >100 kg PASI 75 PASI 90 PASI 100	<b>+ ⊕ +  </b>   <b>+ ⊕ +  </b>   <b>+ ⊕ +  </b>	3.61 (2.80, 4.64)* 3.22 (2.56, 4.03)* 3.39 (2.68, 4.27)*	H-0-1 H-0-1 H-0-1	3.45 (2.67, 4.45)* 3.10 (2.45, 3.89)* 3.27 (2.56, 4.13)*	H#H H#H H#H	3.41 (2.73, 4.34)* 3.07 (2.51, 3.81)* 3.23 (2.62, 4.03)*	<b>→→</b> <b>→</b> <b>→</b>	2.85 (2.16, 3.80)* 2.62 (2.03, 3.40)* 2.77 (2.12, 3.62)*
Adalimumab 40 mg PASI 75 PASI 90 PASI 100	<b>+ ⊕ +  </b>   <b>+ ⊕ +  </b>   <b>+ ⊕ +  </b>	3.64 (2.88, 4.64)* 3.25 (2.63, 4.03)* 3.42 (2.76, 4.26)*	H⊕H H⊕H H⊕H	3.48 (2.67, 4.60)* 3.12 (2.45, 4.01)* 3.29 (2.57, 4.27)*	H <b>⊕</b> -1 H⊕-1 H⊕-1	3.44 (2.70, 4.51)* 3.10 (2.49, 3.95)* 3.27 (2.60, 4.19)*	H <del>0-1</del> H <del>0-1</del> H <del>0-1</del>	2.88 (2.27, 3.70)* 2.65 (2.14, 3.31)* 2.80 (2.24, 3.52)*
Certolizumab pegol 200 mg PASI 75 PASI 90 PASI 100	<b>→ → → → → → → → → →</b>	4.24 (2.94, 6.08)* 3.77 (2.68, 5.31)* 4.04 (2.80, 5.90)*	<b>→</b>	4.05 (2.83, 5.80)* 3.62 (2.58, 5.10)* 3.89 (2.70, 5.69)*	<b>→</b>	4.01 (2.81, 5.78)* 3.59 (2.57, 5.08)* 3.86 (2.69, 5.66)*	<b>→</b>	3.35 (2.31, 4.87)* 3.07 (2.16, 4.37)* 3.31 (2.27, 4.88)*
Tildrakizumab 200 mg PASI 75 PASI 90 PASI 100	<b>⊢●</b> -1 <b>⊢●</b> -1	4.49 (3.23, 6.16)* 3.99 (2.94, 5.37)* 4.32 (3.11, 5.97)*	<b>→</b>	4.29 (3.09, 5.93)* 3.83 (2.82, 5.20)* 4.16 (2.99, 5.80)*	<b>→ → → → → → → → → →</b>	4.24 (3.11, 5.85)* 3.80 (2.84, 5.13)* 4.12 (3.00, 5.72)*	<b>→ → → → → → → → → →</b>	3.54 (2.53, 4.94)* 3.25 (2.37, 4.43)* 3.53 (2.51, 4.95)*
Tildrakizumab 100 mg PASI 75 PASI 90 PASI 100		4.88 (3.53, 6.73)* 4.34 (3.20, 5.86)* 4.76 (3.42, 6.62)*	<b>→</b>	4.67 (3.38, 6.46)* 4.17 (3.08, 5.67)* 4.58 (3.30, 6.42)*	<b>→ → → →</b>	4.62 (3.39, 6.40)* 4.13 (3.09, 5.62)* 4.54 (3.30, 6.36)*	<b>→ → →</b>	3.86 (2.77, 5.40)* 3.53 (2.59, 4.85)* 3.90 (2.78, 5.50)*
Etanercept 25 mg BIW / 50 mg QW PASI 75 PASI 90 PASI 100	- <b>0</b> -1  -0-1  -0-1	12.37 (9.15, 16.77)* 11.55 (8.60, 15.54)* 15.61 (11.17, 21.95)*	H+H	11.83 (8.83, 15.91)* 11.11 (8.33, 14.86)* 15.04 (10.82, 21.03)*	<del></del>	11.70 (8.83, 15.88)* 11.00 (8.33, 14.85)* 14.90 (10.81, 21.00)*	<b>→ → → → → → → → → →</b>	9.78 (7.16, 13.45)* 9.41 (6.93, 12.83)* 12.78 (9.02, 18.18)*
Apremilast 30 mg PASI 75 PASI 90 PASI 100	H <b>⊕</b> H H <b>⊕</b> H	18.61 (14.07, 24.79)* 18.28 (13.86, 24.32)* 27.95 (20.26, 38.95)*	H=H	17.79 (13.30, 24.06)* 17.58 (13.15, 23.75)* 26.91 (19.23, 38.13)*	<b>→ → → →</b>	17.59 (13.45, 23.70)* 17.41 (13.30, 23.38)* 26.67 (19.45, 37.59)*	<b>⊢</b>	14.70 (11.00, 19.93)* 14.89 (11.16, 20.10)* 22.88 (16.40, 32.35)*
Dimethyl fumarate PASI 75 PASI 90 PASI 100	Favoring Favoring	19.70 (12.84, 30.56)* 19.53 (12.26, 31.74)* 30.41 (17.10, 55.95)*	g Favoring		oring Favoring	18.63 (12.22, 29.31)* 18.59 (11.76, 30.64)* 29.01 (16.39, 54.15)*	ring Favoring	15.57 (10.10, 24.42)* 15.90 (9.96, 26.05)* 24.87 (13.93, 46.11)*
	0.3 1.0 3.0 10.0 30.0	- 0.3	1.0 3.0 10.0 30.0	0.3	1.0 3.0 10.0 30.0	0.3	1.0 3.0 10.0 30.0	

BIW, twice a week; kg, kilogram; mg, milligram; NMA, network meta-analysis; PASI, Psoriasis Area and Severity Index; PASI 75, 90, 100, a 75%, 90% or 100% decrease from baseline PASI; QW, once weekly

**eTable 2.** Estimated response rates from the NMA of short-term PASI (sensitivity analyses including global trials only, including phase III trials only, and in an expanded treatment space)

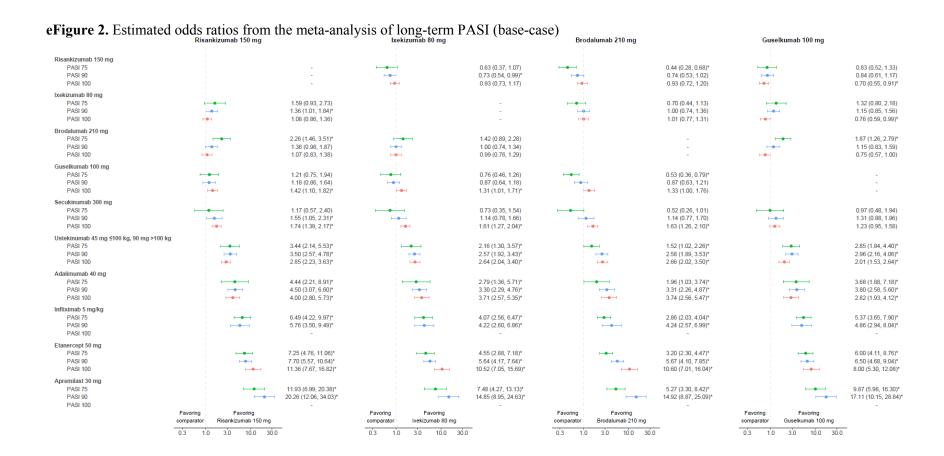
	Sensitivity analysis: global trials only								
Treatment	PA	ASI 75	P	PASI 90	PASI 100				
	Posterior M	ledian, 95% CrI	Posterior N	Median, 95% CrI	Posterior Median, 95% Crl				
Risankizumab 150 mg	89.4%	(87.2%, 91.3%)	71.7%	(67.8%, 75.4%)	41.1%	(36.9%, 45.6%)			
Ixekizumab 80 mg	88.5%	(86.2%, 90.5%)	70.0%	(66.1%, 73.7%)	39.3%	(35.2%, 43.5%)			
Brodalumab 210 mg	88.0%	(85.8%, 90.1%)	69.2%	(65.4%, 73.0%)	38.3%	(34.4%, 42.6%)			
Guselkumab 100 mg	87.3%	(84.4%, 90.0%)	68.0%	(63.2%, 72.9%)	37.1%	(32.2%, 42.6%)			
Secukinumab 300 mg	82.6%	(79.9%, 85.2%)	60.4%	(56.5%, 64.5%)	29.7%	(26.3%, 33.5%)			
Infliximab 5 mg/kg	79.9%	(75.6%, 83.7%)	56.5%	(50.8%, 62.2%)	26.3%	(21.8%, 31.3%)			
Certolizumab pegol 400 mg	71.0%	(65.7%, 75.9%)	45.2%	(39.4%, 51.2%)	18.0%	(14.3%, 22.2%)			
Ustekinumab 45 mg ≤ 100 kg, 90 mg > 100 kg	69.4%	(66.1%, 72.7%)	43.4%	(39.8%, 47.2%)	16.8%	(14.5%, 19.3%)			
Adalimumab 40 mg	70.2%	(66.7%, 73.5%)	44.3%	(40.4%, 48.1%)	17.3%	(14.9%, 20.0%)			
Certolizumab pegol 200 mg	66.1%	(59.9%, 71.9%)	39.8%	(33.6%, 46.3%)	14.6%	(11.1%, 18.7%)			
Tildrakizumab 100 mg	63.0%	(57.8%, 68.1%)	36.6%	(31.7%, 41.9%)	12.7%	(10.1%, 15.8%)			
Tildrakizumab 200 mg	64.9%	(59.8%, 70.0%)	38.6%	(33.5%, 44.0%)	13.8%	(11.1%, 17.2%)			
Etanercept 25 mg BIW / 50 mg QW	39.8%	(35.3%, 44.6%)	17.6%	(14.7%, 20.9%)	4.2%	(3.2%, 5.4%)			
Apremilast 30 mg	31.8%	(27.7%, 36.2%)	12.6%	(10.2%, 15.3%)	2.6%	(1.9%, 3.4%)			
Dimethyl fumarate	30.9%	(23.4%, 39.7%)	12.1%	(8.1%, 17.5%)	2.5%	(1.4%, 4.2%)			

Placebo	5.1%	(4.6%, 5.7%)	1.1%	(0.9%, 1.2%)	0.1%	(0.1%, 0.1%)
	Sens	sitivity analysis: phase	e III trials only	7		
Treatment	PA	ASI 75	P	ASI 90	PA	ASI 100
	Posterior M	ledian, 95% CrI	Posterior M	ledian, 95% CrI	Posterior M	ledian, 95% CrI
Risankizumab 150 mg	89.2%	(86.9%, 91.2%)	71.6%	(67.5%, 75.4%)	40.3%	(35.8%, 44.8%)
Ixekizumab 80 mg	88.7%	(86.4%, 90.7%)	70.6%	(66.7%, 74.4%)	39.2%	(35.0%, 43.5%)
Brodalumab 210 mg	87.6%	(85.1%, 89.8%)	68.7%	(64.6%, 72.7%)	37.1%	(32.9%, 41.5%)
Guselkumab 100 mg	86.8%	(83.9%, 89.5%)	67.4%	(62.6%, 72.0%)	35.7%	(31.0%, 40.8%)
Secukinumab 300 mg	82.8%	(80.0%, 85.4%)	61.0%	(56.9%, 65.1%)	29.5%	(26.0%, 33.3%)
Infliximab 5 mg/kg	80.0%	(75.5%, 84.2%)	57.0%	(51.0%, 63.2%)	26.0%	(21.3%, 31.6%)
Certolizumab pegol 400 mg	69.3%	(62.7%, 75.3%)	43.6%	(36.6%, 50.6%)	16.4%	(12.3%, 21.2%)
Ustekinumab 45 mg ≤ 100 kg, 90 mg > 100 kg	69.1%	(65.6%, 72.4%)	43.3%	(39.6%, 47.2%)	16.2%	(13.9%, 18.8%)
Adalimumab 40 mg	69.9%	(66.3%, 73.3%)	44.2%	(40.3%, 48.3%)	16.8%	(14.3%, 19.5%)
Certolizumab pegol 200 mg	65.3%	(58.5%, 71.6%)	39.2%	(32.6%, 46.2%)	13.8%	(10.2%, 18.0%)
Tildrakizumab 100 mg	62.1%	(56.0%, 68.0%)	36.0%	(30.3%, 42.1%)	12.0%	(9.1%, 15.5%)
Tildrakizumab 200 mg	62.5%	(56.3%, 68.3%)	36.4%	(30.6%, 42.5%)	12.2%	(9.3%, 15.7%)
Etanercept 25 mg BIW / 50 mg QW	40.0%	(35.0%, 45.2%)	17.9%	(14.7%, 21.6%)	4.1%	(3.1%, 5.5%)
Apremilast 30 mg	31.0%	(26.5%, 35.9%)	12.3%	(9.8%, 15.2%)	2.4%	(1.7%, 3.3%)
Dimethyl fumarate	29.8%	(22.1%, 38.6%)	11.6%	(7.6%, 17.0%)	2.2%	(1.2%, 3.8%)

Placebo	5.2%	(4.7%, 5.8%)	1.1%	(0.9%, 1.3%)	0.1%	(0.1%, 0.1%)
	Sensitiv	vity analysis: expande	d treatment sp	pace		
Treatment	P	ASI 75	PA	ASI 90	PA	ASI 100
	Posterior M	ledian, 95% CrI	Posterior M	ledian, 95% CrI	Posterior M	ledian, 95% CrI
Risankizumab 150 mg	89.2%	(86.9%, 91.3%)	71.6%	(67.5%, 75.5%)	40.4%	(35.8%, 45.0%)
Ixekizumab 80 mg	88.4%	(86.0%, 90.5%)	70.1%	(66.0%, 73.9%)	38.7%	(34.3%, 43.0%)
Brodalumab 210 mg	88.7%	(86.4%, 90.9%)	70.7%	(66.8%, 74.8%)	39.4%	(35.2%, 44.1%)
Guselkumab 100 mg	86.8%	(83.7%, 89.4%)	67.3%	(62.5%, 72.0%)	35.7%	(30.9%, 40.8%)
Secukinumab 300 mg	83.5%	(80.8%, 86.1%)	62.1%	(58.0%, 66.2%)	30.6%	(27.0%, 34.6%)
Infliximab 5 mg/kg	80.3%	(76.4%, 83.9%)	57.4%	(52.1%, 62.7%)	26.5%	(22.2%, 31.2%)
Certolizumab pegol 400 mg	71.1%	(65.2%, 76.6%)	45.6%	(39.2%, 52.3%)	17.7%	(13.8%, 22.4%)
Ustekinumab 45 mg ≤ 100 kg, 90 mg > 100 kg	69.8%	(66.4%, 73.3%)	44.1%	(40.3%, 48.2%)	16.8%	(14.4%, 19.5%)
Adalimumab 40 mg	69.6%	(66.2%, 72.8%)	43.9%	(40.2%, 47.6%)	16.6%	(14.3%, 19.1%)
Certolizumab pegol 200 mg	66.2%	(59.4%, 72.6%)	40.2%	(33.4%, 47.4%)	14.4%	(10.6%, 18.9%)
Tildrakizumab 100 mg	63.0%	(57.2%, 68.6%)	36.9%	(31.4%, 42.7%)	12.5%	(9.7%, 15.9%)
Tildrakizumab 200 mg	65.0%	(59.3%, 70.5%)	38.9%	(33.3%, 45.0%)	13.6%	(10.6%, 17.3%)
Etanercept 25 mg BIW / 50 mg QW	40.2%	(35.5%, 45.3%)	18.0%	(14.9%, 21.7%)	4.2%	(3.2%, 5.5%)
Ciclosporin 2.5–3 mg/kg	43.7%	(32.6%, 55.4%)	20.4%	(13.2%, 29.8%)	5.0%	(2.7%, 8.9%)
Methotrexate (high dose)	44.4%	(34.5%, 54.4%)	20.9%	(14.3%, 28.9%)	5.2%	(3.0%, 8.5%)

Methotrexate (low dose)	31.2%	(24.3%, 39.1%)	12.4%	(8.6%, 17.2%)	2.4%	(1.5%, 3.9%)
Apremilast 30 mg	30.9%	(26.8%, 35.1%)	12.2%	(9.9%, 14.7%)	2.4%	(1.8%, 3.1%)
Fumaric acid esters	31.2%	(25.1%, 37.6%)	12.3%	(9.0%, 16.3%)	2.4%	(1.6%, 3.6%)
Dimethyl fumarate	28.0%	(20.7%, 36.2%)	10.6%	(6.9%, 15.4%)	2.0%	(1.1%, 3.3%)
Acitretin 0.4 mg/kg	19.5%	(4.6%, 47.7%)	6.3%	(0.9%, 23.5%)	1.0%	(0.1%, 6.2%)
Placebo	5.4%	(4.8%, 5.9%)	1.1%	(1.0%, 1.3%)	0.1%	(0.1%, 0.1%)

BIW, twice a week; CI, confidence interval; CrI, credible interval; kg, kilogram; mg, milligram; NMA, network meta-analysis; PASI, Psoriasis Area and Severity Index; PASI 75, 90, 100, a 75%, 90% or 100% decrease from baseline PASI; QW, once weekly



BIW, twice a week; kg, kilogram; mg, milligram; PASI, psoriasis area and severity index; PASI 75, 90, 100: a 75%, 90% or 100% decrease from baseline PASI

**eTable 3.** Estimated response rates from the meta-analysis of long-term PASI (sensitivity analyses: including global trials only, including trials reporting NRI data, and including phase III trials only)

	Se	nsitivity analysis: glob	oal trials only			
Treatment	P	ASI 75	P	ASI 90	PA	ASI 100
	Response	Rate, 95% CI	Response	e Rate, 95% CI	Response Rate, 95% CI	
Risankizumab 150 mg	90.1%	(86.3%, 92.9%)	79.4%	(75.5%, 82.9%)	56.2%	(52.4%, 59.9%)
Guselkumab 100 mg	87.8%	(83.8%, 91.0%)	76.3%	(71.4%, 80.6%)	47.4%	(42.1%, 52.8%)
Brodalumab 210 mg	80.0%	(75.7%, 83.7%)	74.0%	(69.3%, 78.1%)	54.5%	(49.5%, 59.4%)
Ixekizumab 80 mg	85.0%	(79.2%, 89.4%)	73.9%	(69.9%, 77.5%)	54.3%	(50.0%, 58.5%)
Secukinumab 300 mg	88.6%	(80.6%, 93.6%)	71.3%	(64.2%, 77.5%)	42.4%	(38.5%, 46.4%)
Ustekinumab 45 mg≤100 kg, 90 mg>100 kg	72.5%	(65.9%, 78.2%)	52.4%	(47.1%, 57.7%)	31.0%	(27.2%, 35.2%)
Adalimumab 40 mg	67.1%	(52.9%, 78.7%)	46.2%	(38.6%, 53.9%)	24.2%	(18.8%, 30.7%)
Infliximab 5 mg/kg	58.3%	(52.5%, 63.8%)	40.1%	(30.0%, 51.1%)	-	-
Etanercept 50 mg	55.5%	(50.1%, 60.9%)	33.4%	(28.5%, 38.7%)	10.1%	(7.3%, 13.9%)
Apremilast 30 mg	44.7%	(29.4%, 61.0%)	16.0%	(10.7%, 23.3%)	-	-
	Sensiti	vity analysis: trials re	porting NRI d	lata		
Treatment	P	ASI 75	P	ASI 90	PA	ASI 100
	Response	Rate, 95% CI	Response	Rate, 95% CI	Response	Rate, 95% CI
Risankizumab 150 mg	90.1%	(86.3%, 92.9%)	79.4%	(75.5%, 82.9%)	56.2%	(52.4%, 59.9%)

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(88.1%, 94.1%) (75.7%, 83.7%) (79.2%, 89.4%)	74.9% 74.0% 73.9%	(69.9%, 79.2%) (69.3%, 78.1%)	44.9% 54.5%	(39.7%, 50.3%) (49.5%, 59.4%)
, , ,		` , , ,	54.5%	(49.5%, 59.4%)
(79.2%, 89.4%)	73.9%	(60.00/ 77.50/)		
		(69.9%, 77.5%)	54.3%	(50.0%, 58.5%)
(64.2%, 77.0%)	52.4%	(47.1%, 57.7%)	31.0%	(27.2%, 35.2%)
(56.8%, 66.5%)	41.9%	(28.5%, 56.7%)	22.0%	(16.7%, 28.4%)
(52.5%, 63.8%)	40.1%	(30.0%, 51.1%)	-	-
(50.1%, 60.9%)	33.4%	(28.5%, 38.7%)	10.1%	(7.3%, 13.9%)
(30.2%, 50.7%)	-	-	-	-
	(56.8%, 66.5%) (52.5%, 63.8%) (50.1%, 60.9%)	(56.8%, 66.5%) 41.9% (52.5%, 63.8%) 40.1% (50.1%, 60.9%) 33.4%	(56.8%, 66.5%)       41.9%       (28.5%, 56.7%)         (52.5%, 63.8%)       40.1%       (30.0%, 51.1%)         (50.1%, 60.9%)       33.4%       (28.5%, 38.7%)	(56.8%, 66.5%)       41.9%       (28.5%, 56.7%)       22.0%         (52.5%, 63.8%)       40.1%       (30.0%, 51.1%)       -         (50.1%, 60.9%)       33.4%       (28.5%, 38.7%)       10.1%

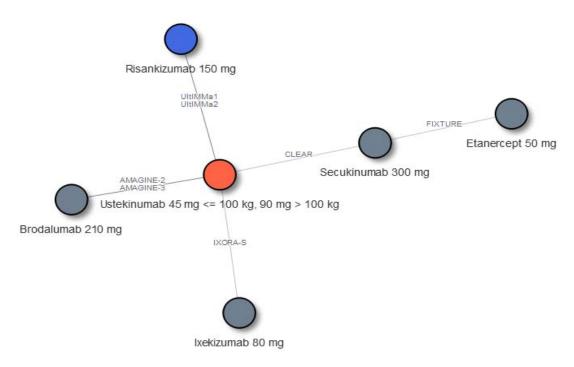
## Sensitivity analysis: phase III trials only

Treatment	PASI 75 Response Rate, 95% CI		PASI 90 Response Rate, 95% CI		PASI 100 Response Rate, 95% CI	
Risankizumab 150 mg	90.1%	(86.3%, 92.9%)	79.4%	(75.5%, 82.9%)	56.2%	(52.4%, 59.9%)
Guselkumab 100 mg	88.2%	(84.6%, 91.1%)	76.5%	(72.1%, 80.5%)	47.4%	(42.5%, 52.4%)
Brodalumab 210 mg	80.0%	(75.7%, 83.7%)	74.0%	(69.3%, 78.1%)	54.5%	(49.5%, 59.4%)
Ixekizumab 80 mg	85.0%	(79.2%, 89.4%)	73.9%	(69.9%, 77.5%)	54.3%	(50.0%, 58.5%)
Secukinumab 300 mg	88.6%	(80.6%, 93.6%)	71.3%	(64.2%, 77.5%)	42.4%	(38.5%, 46.4%)
Ustekinumab 45 mg ≤ 100 kg, 90 mg > 100 kg	72.5%	(65.9%, 78.2%)	52.4%	(47.1%, 57.7%)	31.0%	(27.2%, 35.2%)
Adalimumab 40 mg	71.4%	(52.1%, 85.1%)	48.7%	(44.1%, 53.3%)	25.6%	(20.2%, 31.9%)

Infliximab 5 mg/kg	58.3%	(52.5%, 63.8%)	40.1%	(30.0%, 51.1%)	-	-
Etanercept 50 mg	55.5%	(50.1%, 60.9%)	33.4%	(28.5%, 38.7%)	10.1%	(7.3%, 13.9%)
Apremilast 30 mg	52.7%	(41.4%, 63.8%)	17.6%	(10.5%, 27.9%)	-	-

BIW, twice a week; CI: confidence interval; CrI, credible interval; kg, kilogram; mg, milligram; NRI, non-response imputation; PASI, Psoriasis Area and Severity Index; PASI 75, 90, 100, a 75%, 90% or 100% decrease from baseline PASI

eFigure 3. Evidence network for NMA of long-term PASI



kg, kilogram; mg, milligram; NMA, network meta-analysis; PASI, Psoriasis Area and Severity Index

eTable 4. Estimated response rates from the NMA of long-term PASI

Treatment	PASI 75		PASI 90		PASI 100	
	Posterior Median, 95% CrI		Posterior Median, 95% CrI		Posterior Median, 95% CrI	
Risankizumab 150 mg	91.1%	(87.6%, 93.8%)	81.3%	(75.7%, 86.1%)	59.7%	(52.1%, 67.1%)
Brodalumab 210 mg	88.2%	(84.7%, 91.1%)	76.6%	(71.3%, 81.4%)	53.3%	(46.7%, 59.9%)
Ixekizumab 80 mg	83.5%	(76.2%, 89.2%)	69.8%	(60.1%, 78.3%)	45.0%	(34.8%, 55.6%)
Secukinumab 300 mg	80.1%	(74.7%, 84.8%)	65.2%	(58.1%, 71.6%)	39.9%	(32.9%, 47.3%)
Ustekinumab 45 mg≤100 kg, 90 mg>100 kg	69.8%	(67.3%, 72.3%)	52.5%	(49.3%, 55.7%)	28.0%	(25.2%, 31.0%)
Etanercept 50 mg BIW	53.8%	(44.1%, 63.5%)	35.9%	(27.1%, 45.6%)	15.7%	(10.4%, 22.6%)

CrI, credible interval; kg, kilogram; mg, milligram; NMA, network meta-analysis; PASI, Psoriasis Area and Severity Index; PASI 75, 90, 100, a 75%, 90% or 100% decrease from baseline PASI

eTable 5. Estimated Odds Ratios for PASI 90 from the NMA of long-term PASI

Etanercept 50 mg	1.97 (1.35, 2.91)*	3.34 (2.53, 4.42)*	4.13 (2.35, 7.34)*	5.86 (3.72, 9.36)*	7.80 (4.75, 12.94)*
0.51 (0.24 0.74)*	11.11.1.1.1.1.1.1.1.1.1.1.1.1.1.1.1.1.1.	1.60 (1.20, 2.22)*	2.00 (1.20.2.21)*	2.07 (2.22. 2.02)*	2.05 (2.00, 5.45)*
0.51 (0.34, 0.74)*	Ustekinumab 45 mg ≤100 kg, 90 mg >100 kg	1.69 (1.29, 2.22)*	2.09 (1.39, 3.21)*	2.97 (2.32, 3.83)*	3.95 (2.90, 5.45)*
0.30 (0.23, 0.39)*	0.59 (0.45, 0.78)*	Secukinumab 300 mg	1.24 (0.75, 2.05)	1.76 (1.22, 2.54)*	2.33 (1.54, 3.56)*
0.24 (0.14, 0.42)*	0.48 (0.31, 0.72)*	0.81 (0.49, 1.32)	Ixekizumab 160 mg	1.42 (0.87, 2.30)	1.89 (1.12, 3.19)*
0.17 (0.11, 0.27)*	0.34 (0.26, 0.43)*	0.57 (0.39, 0.82)*	0.70 (0.44, 1.15)	Brodalumab 210 mg	1.33 (0.89, 1.99)
0.13 (0.08, 0.21)*	0.25 (0.18, 0.34)*	0.43 (0.28, 0.65)*	0.53 (0.31, 0.90)*	0.75 (0.50, 1.12)	Risankizumab 150
					mg